## S5 Appendix. STROBE Statement

	Item No	Recommendation	Location
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	Title
		(b) Provide in the abstract an informative and balanced summary of	Abstract,
		what was done and what was found	paragraph 2
Introduction			hama-Arahm =
Background/rationale	2	Explain the scientific background and rationale for the investigation	Introduction,
	_	being reported	paragraph 2
Objectives	3	State specific objectives, including any prespecified hypotheses	Introduction,
		3 , 1 ,	paragraph 2
Methods			
Study design	4	Present key elements of study design early in the paper	Introduction,
			paragraph
			2; Methods,
			paragraph 1
Setting	5	Describe the setting, locations, and relevant dates, including periods of	Methods
		recruitment, exposure, follow-up, and data collection	paragraph 1
Participants	6	(a) Give the eligibility criteria, and the sources and methods of selection	Methods
·		of participants	paragraph 1
Variables	7	Clearly define all outcomes, exposures, predictors, potential	Methods,
		confounders, and effect modifiers. Give diagnostic criteria, if applicable	paragraphs
			5-6
Data sources/	8*	For each variable of interest, give sources of data and details of	Methods,
measurement		methods of assessment (measurement). Describe comparability of	paragraphs
		assessment methods if there is more than one group	2-4
Bias	9	Describe any efforts to address potential sources of bias	Methods,
			paragraphs
			5-6
Study size	10	Explain how the study size was arrived at	n/a
Quantitative	11	Explain how quantitative variables were handled in the analyses. If	Methods,
variables		applicable, describe which groupings were chosen and why	paragraphs
			5-6
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	Methods,
		confounding	paragraphs
			5-6
		(b) Describe any methods used to examine subgroups and interactions	Methods,
			paragraph 5
		(c) Explain how missing data were addressed	n/a
		(d) If applicable, describe analytical methods taking account of	n/a
		sampling strategy	
		(e) Describe any sensitivity analyses	n/a
Results			
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers	Results,
		potentially eligible, examined for eligibility, confirmed eligible, included	paragraph 1
		in the study, completing follow-up, and analysed	

		(b) Give reasons for non-participation at each stage	n/a
		(c) Consider use of a flow diagram	n/a
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical,	Table 1
		social) and information on exposures and potential confounders	
		(b) Indicate number of participants with missing data for each variable	n/a
		of interest	
Outcome data	15*	Report numbers of outcome events or summary measures	Table S3 in
			S3
			Appendix
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	Unadjusted
		estimates and their precision (eg, 95% confidence interval). Make clear	estimates:
		which confounders were adjusted for and why they were included	S3
			Appendix;
			Adjusted
			estimates:
			Figs 1-2
		(b) Report category boundaries when continuous variables were	Methods,
		categorized	paragraph 5
		(c) If relevant, consider translating estimates of relative risk into	n/a
		absolute risk for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and	S4
		interactions, and sensitivity analyses	Appendix
Discussion			
Key results	18	Summarise key results with reference to study objectives	Discussion,
			paragraph 1
Limitations	19	Discuss limitations of the study, taking into account sources of potential	Discussion,
		bias or imprecision. Discuss both direction and magnitude of any	paragraph 5
		potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	Discussion,
		limitations, multiplicity of analyses, results from similar studies, and	paragraphs
		other relevant evidence	2-3
Generalisability	21	Discuss the generalisability (external validity) of the study results	Discussion,
			paragraphs
			3-4
Other information			
Funding	22	Give the source of funding and the role of the funders for the present	Provided to
		study and, if applicable, for the original study on which the present	journal
		article is based	

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.